PEPTIDE MIMICS OF CONSERVED GONOCOCCAL EPITOPES AND METHODS AND COMPOSITIONS USING THEM

TECHNICAL FIELD OF THE INVENTION

5   The present invention relates to peptide mimics of conserved epitopes of *Neisseria gonorrhoeae*, which epitopes are not found on human blood group antigens. This invention also relates to methods and compositions using such peptide mimics for the

10   prophylaxis of gonorrheal infections.

BACKGROUND OF THE INVENTION

The sexually transmitted disease, gonorrhea, poses a worldwide risk as one of the most commonly reported communicable diseases. Gonorrhea is caused by

15   the bacterium *Neisseria gonorrhoeae*, a gram negative diplococcus. Although the pathogen primarily infects mucous membranes, it is capable of invading tissues and evading host defenses. *N. gonorrhoeae* is the causative agent of a spectrum of sequelae. These range from

20   asymptomatic mucosal infection to significant disease syndromes in both men and women. The more serious of such syndromes include, for example, disseminated gonococcal infection ("DGI") in men and women, as well as salpingitis or pelvic inflammatory disease ("PID")

25   in women. Either salpingitis or PID may themselves

lead to long-term sequelae, including ectopic pregnancy and infertility. Other important sequelae, sometimes requiring surgical intervention, include recurrent infection, chronic pelvic pain, dyspareunia, pelvic
5 adhesions and other inflammatory residua.

     It has been estimated that in the United States, the direct and indirect costs of treating PID and associated ectopic pregnancy and infertility totaled 2.6 billion dollars in 1984 (53). The total
10 direct costs were estimated to be 2.18 billion dollars in 1990, with indirect costs of 1.54 billion dollars. Assuming constant inflation and incidence rates of PID, the total cost of this disease is projected to reach 8 billion dollars in the year 2000 (9).

15   Despite public health efforts to control gonococcal infections and the availability of effective antibiotic therapies in the United States, there are approximately 315,000 cases of gonorrhea reported annually to the Centers for Disease Control ("CDC")
20 (12). A substantial proportion of all cases of gonorrhea occur in asymptomatically infected individuals who are the source of most new cases within a community (6). The increasing prevalence of antibiotic-resistant strains has complicated treatment
25 of the infection (10, 11, 52).

     N. gonorrhoeae has multiple virulence factors. The surface components of this pathogen play an important role in attaching to and invading host cells, while providing potential targets for the host
30 immune response. Gonococcal infections elicit local and systemic humoral and cellular immune responses to several components which are exhibited as surface exposed antigens of the bacterium, particularly pili, porin ("Por") or protein I ("PI"), opacity associated

proteins ("Opas") or protein IIs, Rmp or protein III, and lipooligosaccharides ("LOSs") (7). Pili, Opa, Por and LOS are all implicated in attachment to and invasion of the host and all display considerable variation on their surface exposed regions (26, 45, 46). The intra-and inter-strain variations of gonococcal surface components have led to hypotheses regarding tissue specificity at different sites and the organism's potential for reinfection and continued virulence.

In both symptomatic and asymptomatic patients, gonococcal infections have been shown to stimulate increased levels of anti-gonococcal serum immunoglobulins. The peripheral humoral response is predominately IgG (mostly subclass IgG3), with lesser amounts of IgM and IgA (13). Quantitatively, the antibody response is primarily directed against the pili, Opa proteins and LOS. Local antibodies are present in genital secretions, but in reduced amounts (48), and may be directed against different antigenic targets than those in serum (27). The predominant class of antibodies present in secretions is also IgG (mostly IgG3) and not secretory IgA ("sIgA") (7). Antibodies against LOS are present as well, but in lesser amounts than those against pili, Por and Opa. Although patients infected with *N. gonorrhoeae* may show an antibody response to many gonococcal antigens, *N. gonorrhoeae* isolated from patients with disseminated infection (DGI) are resistant to the bactericidal action of normal human serum ("NHS") and of most convalescent sera (38). This serum-resistant phenotype, termed stable serum resistance ("SR"), may enable the organism to evade local defenses, penetrate mucosal barriers and disseminate via the bloodstream.

Upon subculture, many strains of gonococci become phenotypically sensitive to killing by NHS or serum sensitive (38). These organisms are termed serum sensitive ("SS") or unstably serum-resistant. Such organisms are frequently isolated from women with severe manifestations of local inflammation or clinically evident PID. Acute salpingitis, the pathologic counterpart of PID (caused by SS gonococci), rarely progresses to bacteremic illness or DGI. This suggests that the intense local inflammatory response, generated by SS gonococci, may serve to contain the infection and prevent bacteremia, although at the cost of damaging the local tissues. SS gonococci generate significantly greater amounts of the complement derived chemotactic peptide, C5a, than do SR gonococci (16). This may be responsible for the polymorphonuclear leukocyte ("PMN") mediated inflammatory response that is produced by SS gonococci.

The development of antibiotic-resistant strains of *N. gonorrhoeae*, has rendered control of this infection increasingly difficult. The potential to undertreat gonococcal infection has accelerated the need for an anti-gonococcal vaccine. The prevention of gonococcal infection, particularly the severe complications of PID, has been the goal of many investigators. Ongoing attempts to develop an effective anti-gonococcal vaccine, however, have been plagued with several difficulties.

Attempts to use individual surface components of the pathogen as targets for conventional vaccines have been unsuccessful because of their antigenic variability. Pilus vaccines have been protective only against infection with the homologous strain (used to make the pilus vaccine) and Por vaccination has been

unsuccessful even in human experimental challenge. In
addition, *N. gonorrhoeae* express marked phenotypic
heterogeneity, typically shifting from one antigenic
form to another at a frequency of >1 in 10^3 organisms
(49, 50) making the surface of this organism a moving
target for most vaccine strategies. Although the
vaccine candidates have provoked antibody responses,
the antibodies and immune responses produced have not
been broadly protective.

LOS is an important virulence determinant of
N. gonorrhoeae. Considerable evidence supports the
role of LOS as a major target of bactericidal antibody
directed to the surface of *N. gonorrhoeae* (2, 16, 18,
37, 47). Antibodies to LOS have several important
functions: bactericidal activity, complement
activation through the classical or alternative
complement pathways (2), and opsonic activity (16).
Additionally, LOS has been shown to be the most
effective gonococcal antigen to induce a functional
antibody response to homologous and heterologous
gonococci (51).

The monoclonal antibody ("mAb") 2C7 (30),
detects a LOS derived oligosaccharide ("OS") epitope
that appears to be widely conserved and expressed
amongst clinical isolates of gonococci. Typically,
saccharides are T-cell independent antigens. When
administered alone as immunogens, they generally elicit
only a primary antibody response. In addition,
oligosaccharides are small (<10 saccharide units) (19),
and would likely require additional biochemical
derivatization to render them immunogenic. The use of
such oligosaccharides as vaccine candidates, therefore,
is limited in several respects.

Internal image determinants have been proposed for use in vaccines (36). By means of mAb technology, a protective antibody (Ab1) to an epitope of interest on the pathogen can be produced. The

5 particular antibody (Ab1) can be purified and subsequently used as an immunogen to elicit an anti-idiotypic antibody (Ab2) which may be an internal image of the original epitope on the pathogen.

As predicted by the Jerne "network" theory

10 (23), immunization with an anti-idiotypic antibody (Ab2) that is directed against antigen combining sites of primary antibody (Ab1), may elicit a humoral immune response specific for the nominal antigen. The resulting anti-anti-idiotypic antibody (or Ab3) should

15 react with the original primary antigen. If the primary antigen is an oligosaccharide (and therefore expected to give a T-cell independent immune response), then immunization with Ab2 (the protein equivalent) may elicit a T-cell dependent response.

20 It has been demonstrated that an anti-idiotope of mAb 2C7 elicits anti-LOS antibodies in mice and rabbits that together with complement are bactericidal for gonococci, and that serum from animals immunized with this anti-idiotypic antibody also

25 supports opsonophagocytosis by human PMNs (20).

It has also been shown that synthetic peptides which mimic a nominal antigen through binding to a specific antibody directed to the nominal antigen can also elicit an immune response against the nominal

30 antigen (29, 24, 54).

The need exists for an agent useful for the prevention of gonorrhea targeted to the prevention of gonococcal salpingitis, an infection that may be associated with debilitating and chronic pelvic pain,

infertility and ectopic pregnancy (42). Another important objective is to prevent transmission of the organism from an infected but asymptomatic host to an otherwise immune sexual consort. This is important
5 because a substantial fraction of all cases of gonorrhea in both men and women are asymptomatic, and asymptomatically infected, sexually active persons are probably the major source of most new infections. Accordingly, a gonococcal vaccine that only attenuates
10 the severity of symptomatic gonorrhea could result in a higher ratio of asymptomatic/symptomatic cases and as a result, such a vaccine might promote the spread of gonorrhea, unless it also prevents transmission (41).

SUMMARY OF THE INVENTION

15 The present invention generally solves the problems referred to above by providing peptide mimics of widely conserved oligosaccharide epitopes of *N. gonorrhoeae* which are not present in human blood group antigens. Also provided are methods for
20 producing the peptide mimics according to this invention.

The peptide mimics according to this invention are useful in methods and compositions for the prophylaxis of *N. gonorrhoeae* infections.

25 ## BRIEF DESCRIPTION OF THE DRAWINGS

Figure 1 shows a Western blot analysis of the binding of mAb 2C7 to *E. coli* clones. The seven unique *E. coli* clones (PEP1-PEP7) [SEQ ID NOS:1-7] were grown in IMC media containing 100 μg/ml ampicillin, and then
30 induced to express fusion proteins. Bacterial lysates from each of the clones were prepared and loaded onto 14% SDS-PAGE gels. After electrophoresis, the proteins

were transferred to Immobilon PVDF transfer membranes using a Biorad electrophoretic transfer apparatus (Biorad, Hercules CA). The membranes were probed with mAb 2C7 (A) or anti-thioredoxin antibody (B). A
5 negative clone that did not bind mAb 2C7 was used as a control [SEQ ID NO:9].

Figure 2 shows the peptide mimic sequences derived from the seven *E.coli* clones that bind to mAb 2C7.

10 Figure 3 shows FACS analysis of mAb 2C7 binding to *E.coli* clones expressing peptide mimic fusions. *E. coli* clones were grown in IMC media containing 100 μg/ml ampicillin, and then induced to express fusion proteins. The bacterial cells were
15 fixed with 1% paraformaldehyde before staining with mAb 2C7, followed by FITC-conjugated anti-mouse IgG. A negative clone that did not bind mAb 2C7 was used as a control [SEQ ID NO:9]. The number below the *E. coli* clones represents the median fluorescent intensity in
20 the populations that bind to mAb 2C7 compared to the control; the number in parenthesis shows the percentage of the cells in the population (total population = 100%).

Figure 4 shows inhibition of mAb 2C7 binding
25 to LOS by *E. coli* clones expressing peptide fusions. *E. coli* clones were grown in IMC media containing 100 μg/ml ampicillin, and then induced to express fusion proteins. *E. coli* cells were incubated with mAb 2C7 for 30 min. before loading onto LOS coated plates. A
30 negative clone that did not bind mAb 2C7 was used as a control [SEQ ID NO:9]. The data represent means from at least 2 experiments (duplicate wells). PEP1 clones showed the maximum inhibition of mAb 2C7 binding to LOS (66%) [SEQ ID NO:1]. PEP7, PEP3, PEP4, PEP2, PEP6, and

PEP5 showed respective decreases in inhibition of
binding [SEQ ID NOS:7, 3, 4, 2, 6 and 5, respectively].

Figure 5 shows inhibition of mAb 2C7 binding
to LOS by a peptide comprising the consensus sequence
5   (DE_GLF) [SEQ ID NO:8]. The data represent means ±SE
from 3 experiments (duplicate wells). Peptide PEP1
inhibited the binding of mAb 2C7 to LOS in a dose
responsive manner.

Figure 6 shows binding of mAb 2C7 to the
10   multiple antigen peptide ("MAP") MAP1.

Figure 7 shows inhibition of mAb 2C7 binding
to LOS by multiple antigen peptides.

Figure 8 shows octa-MAP1-induced IgG anti-LOS
antibody responses in mice. (A) Eight mice received a
15   dose of 50 μg of Octa-MAP1 emulsified in Freund's
adjuvant on day 0 and again on day 21. (B) Four mice
were immunized with purified LOS as a positive control.
Mice were immunized with either Freund's adjuvant (C)
or an unrelated octa-MAP control peptide (D) as
20   negative controls.

Figure 9 shows IgG anti-LOS antibody
responses in all immunized mice. IgG anti-LOS antibody
responses (mean ± SE) are shown for all mice (including
animals that exhibited no response).

25   Figure 10 shows IgG anti-LOS antibody
responses in responder mice only. Antibody response
was defined as IgG anti-LOS (mean ± SE) greater than
0.4 μg/ml (4 fold above baseline IgG anti-LOS levels).
Mice were immunized with Octa-MAP1, LOS, Freund's
30   adjuvant alone or unrelated octa-MAP control peptide.
Elicited IgG anti-LOS antibody levels were plotted as a
function of concentration over time.

Figure 11 shows IgM anti-LOS antibody
responses in responder mice only. Mice were immunized

with Octa-MAP1, LOS, Freund's adjuvant alone or
unrelated octa-MAP control peptide. Elicited IgG anti-
LOS antibody levels were plotted as a function of
concentration over time.

5   Figure 12 shows survival of *gonorrhoeae*
strain 15253 and its *lgtG* mutant (2C7 epitope negative)
exposed to mouse immune serum (67% [100 µl of serum in
150 µl total reaction volume] plus added human
complement from normal human donor serum [giving a
10   final human complement concentration of 17% by
volume]). A bactericidal assay was performed using (A)
mAb 2C7 mice against strain 15253 (positive control)
and strain 15253 *lgtG* (negative control) (4). 25 µg/ml
of mAb 2C7 (100 µl in 150 µl of total volume of
15   reaction mixture) mediated 100% killing of strain
15253, and no killing of strain 15253 *lgtG*. (B) Normal
mouse serum (pool of 20 mouse sera, mean concentration
of IgG anti-LOS antibody, 0.1 µg/ml) failed to kill
either strain. (C) Serum taken from a single mouse
20   immunized with Octa-MAP1 (containing 5.05 µg/ml of IgG
anti-LOS antibody, pooled from bleeds taken between
weeks 7-11) showed 92% killing (8% survival) of strain
15253, whereas strain 15253 *lgtG* survived fully. (D)
Serum taken from a single mouse immunized with LOS
25   (containing 21.98 µg/ml of IgG anti-LOS antibodies,
pooled from bleeds taken between weeks 7-11) showed no
killing of strain 15253 (179% survival) and strain
15253 *lgtG* (133% survival). Single mice immunized with
negative control antigens (E) Freund's adjuvant alone
30   or (F) unrelated octa-MAP control peptide did not kill
either strain. Figure 12 controls included the
Complement source without antibody (137.9% ± 1.0%

survival (no killing) for strain 15253, and 132.5% ±
14.3% survival (no killing) for the *lgtG* mutant of
15253).

    Figure 13 shows a plot of IgG anti-LOS
5 antibody concentration versus killing of *N. gonorrhoeae*
strain 15253. IgG anti-LOS antibody levels from each
of three mice immunized with Octa-MAP1 are plotted
versus percent bacterial killing. Mouse sera
containing 1.38, 2.50 and 5.05 μg/ml of anti-LOS
10 antibodies showed 31, 74 and 92 % killing respectively
of strain 15253. Killing by mAb 2C7 is shown at 5
separate LOS antibody concentrations as a positive
control.

DETAILED DESCRIPTION OF THE INVENTION

15 Definitions

    As used herein, an "antibody" is an intact
immunoglobulin molecule comprising two each of
immunoglobulin light and heavy chains. Accordingly,
antibodies include intact immunoglobulins of types IgA,
20 IgG, IgE, IgD, IgM (as well as subtypes thereof),
wherein the light chains of the immunoglobulin may be
of types kappa or lambda.

    As used herein, "monoclonal antibodies" are
monospecific antibodies produced initially by a single
25 clone of antibody forming cells.

    As used herein, "immunoprophylactically
effective" means the ability to induce in a normal
individual an immune response sufficient to protect
said patient for some period of time against
30 *N. gonorrhoeae* infection.

As used herein, "peptide" means a linear or cyclic chain of amino acids, usually at least 4 and less than 50 amino acids in length.

As used herein, "peptide mimic" means a
5 peptide which exhibits an immunological antibody binding profile similar to that of a known epitope.

PEPTIDE MIMICS AND THEIR USE IN
COMPOSITIONS AND METHODS ACCORDING TO THIS INVENTION

The present invention is directed to peptide
10 mimics that immunospecifically react with an antibody directed to a conserved oligosaccharide epitope of *N. gonorrhoeae*, which oligosaccharide epitope is not present in human blood group antigens. Such peptide mimics can be used in a manner similar to the anti-
15 idiotypic antibodies described, for example in United States patents 5,476,784 and 6,099,839 (both incorporated herein by reference), as a surrogate antigen to elicit a T cell-dependent immune response against an oligosaccharide epitope of *N. gonorrhoeae*.
20 The peptide mimic may be administered to uninfected individuals to induce a specific immune response directed against gonococcal organisms or cells bearing said oligosaccharide antigen. Such an immune response can be immunoprophylactic in character, in
25 that it would prevent an infection should the recipient be exposed to the gonococcal organism or cells bearing said oligosaccharide antigen.

A random peptide library may be screened based on antibody binding specificity in order to
30 identify candidate peptide mimics. The technology for such screening is known to those of skill in the art. In one approach, a random peptide library expressed on *E. coli* flagella may be used to identify peptides that

bind to a conserved oligosaccharide epitope of
N. gonorrhoeae, which oligosaccharide epitope is not
present in human blood group antigens. For example,
binding to mAb 2C7 may be assayed to identify candidate

5 peptide mimics. Binding may be characterized by
western blotting, flow cytometric analysis or
competition for binding of mAb 2C7 to LOS by solid-
phase ELISA.

Antibody modeling may also be used to define

10 an immunogenic site in the complementarity determining
regions (CDRs) of an anti-idiotope corresponding to the
epitope of interest. Such analysis may yield
information about the three-dimensional conformation of
the immunogenic site that is useful in the design of a

15 peptide mimic of the immunogenic site.

Once a specific peptide mimic is identified
and sequenced, it may be produced synthetically by
methods known in the art.

Peptide mimics may also be modified to elicit

20 a greater immune response through the use of haptens,
the use of adjuvants, linking the peptide mimic to a
carrier protein, using a multiple antigen peptide,
coupling the peptide mimic to a complement protein or
through other methods known in the art.

25 The preferred pharmaceutical compositions of
this invention are similar to those used for
immunization of humans with other peptides. Typically,
the peptide mimics of the present invention will be
suspended in a sterile saline solution for therapeutic

30 uses. The pharmaceutical compositions may
alternatively be formulated to control release of the
active ingredients or to prolong their presence in a
patient's system. Numerous suitable drug delivery
systems are known and include, e.g., implantable drug

release systems, hydrogels, hydroxymethylcellulose, microcapsules, liposomes, microemulsions, microspheres, and the like.

The pharmaceutical compositions of this invention may be administered by any suitable means such as orally, intranasally, subcutaneously, intramuscularly, intravenously, intra-arterially, or parenterally. Ordinarily, intravenous (i.v.) or parenteral administration will be preferred.

It will be apparent to those of ordinary skill in the art that the immunoprophylactically effective amount of peptide mimics of this invention will depend, *inter alia*, upon the administration schedule, the unit dose of peptide mimic administered, whether the peptide mimic is administered in combination with other therapeutic agents, the immune status and health of the patient, the therapeutic activity of the peptide mimic administered and the judgment of the treating physician.

In order that this invention may be better understood, the following examples are set forth. These examples are for purposes of illustration only, and are not to be construed as limiting the scope of the invention in any manner.

<u>EXAMPLES</u>

I.   Identification of Clones that Encode
      <u>Peptides that Specifically Bind to mAb 2C7</u>

      A.   <u>Random peptide display</u>
            A FliTrx™ random peptide library (Invitrogen, Carlsbad CA) was used to express peptides (12-mers) of random sequence on the surface of *E. coli*. The DNA encoding this library of peptides is inserted within

the gene encoding the active loop of thioredoxin which is itself inserted into the nonessential region of the flagellin gene. Expression of the peptide fusion is controlled by the bacteriophage lambda major leftward

5 promoter (P_L) in the vector pFliTrx™. In this system, P_L is induced by the addition of tryptophan. When induced, the fusion protein is exported and assembled into flagella on the bacterial cell surface, allowing for the display of the peptide.

10   B.   Screening of peptides that bind to mAb 2C7
               The FliTrx™ peptide library (1.77×10^8
primary clones) was grown overnight in IMC medium (0.2% w/v casamino acid, 0.5% w/v glucose, 42 mM Na_2HPO_4, 22 mM KH_2PO_4, 8.5 mM NaCl, 18.7 mM NH_4Cl and 1mM $MgCl_2$)
15 containing 100 μg/ml ampicillin, at 25°C. The expression of fusion peptides was induced by adding L-tryptophan to a final concentration of 100 μg/ml, and the culture was grown at 25°C for 6 h. The induced peptide fusion library was then incubated with a 2C7
20 mAb-coated plate (20 μg/ml). After 1 h incubation, the plate was washed 5 times with IMC medium containing 100 μg/ml ampicillin and 1% α-methyl mannoside. Bound *E. coli* were eluted by mechanical shearing or by competition with purified LOS prepared from gonococcal
25 strain 15253 (the mAb 2C7 epitope is known to be expressed in strain 15253), and then grown overnight at 25°C. After the fifth round of panning, bound *E. coli* were eluted and plated on RMG agar (2% w/v casamino acid, 0.5% w/v glucose, 42 mM Na_2HPO_4, 22 mM KH_2PO_4, 8.5
30 mM NaCl, 18.7 mM NH_4CI, 1 mM $MgCl_2$, and 1.5% agar) containing 100 μg/ml ampicillin at 25°C. Individual bacterial colonies were chosen to assay binding to mAb 2C7 by western blot (a hybridoma cell line secreting

mAb 2C7 is deposited with the American Type Culture
Collection ["ATCC"] and is assigned ATCC accession
number HB-11859).

5   The library was subjected to 5 rounds of
positive selection with mAb 2C7 coated on a 60 mm
tissue culture plate or subjected to negative selection
for 1 h with irrelevant IgG3 (Sigma, St. Louis, MO)
first, before proceeding to 5 rounds of positive
selection with mAb 2C7.

10   One hundred-seven colonies were randomly
chosen and screened by western blot for the ability to
bind mAb 2C7. Fourteen clones were identified that
bound to mAb 2C7. Plasmid DNA was then prepared from
the positive clones and sequenced using primers that
15   bind to regions that are located at the 5' and 3'
flanks of the inserted peptide's nucleotide sequence.
Seven unique clones were identified, as shown in
Figures 1 and 2 [SEQ ID NOS:1-7].

C.   Flow cytometric analyses

20   Positive *E. coli* clones were grown overnight
in IMC media containing 100 μg/ml ampicillin, at 25°C
and then induced to express the peptide fusions for 6
h. *E. coli* cells were fixed with 0.5% paraformaldehyde
on ice for 10 min. Aliquots of 200-μl of fixed
25   organisms were spun at 2000 x g for 10 min.
Supernatants were discarded, and pellets were
resuspended in blocking buffer (IMC media containing
100 μg/ml ampicillin, 1% nonfat dry milk, 150 mM NaCl
and 1% α-methyl mannoside) containing mAb 2C7.
30   Suspensions were incubated at 37°C for 30 min before
spinning at 2000 x g for 10 min. Pellets were washed
with 100 μl of washing buffer (IMC media containing
100μg/ml ampicillin and 1% α-methyl mannoside) and then

resuspended in 100 μl of blocking buffer containing
FITC-conjugated anti-mouse IgG (Sigma, St. Louis, MO).
The mixtures were incubated at 37°C for 30 min before
spinning at 2000 x g for 10 min. Supernatants were
5   removed, and pellets washed in 100 μl of washing buffer
before resuspension in 1 ml of PBS. The suspensions
were analyzed on a FACS using CellQuest software
(Becton Dickinson, Franklin Lakes NJ). A negative
clone that did not bind mAb 2C7 was used as a control.

10   The binding of *E. coli* cells to mAb 2C7 was
observed to increase from *E. coli* clone PEP3, PEP4,
PEP6, PEP5, PEP2, PEP7 to PEP 1 (according to median
fluorescent intensity, "MFI") [SEQ ID NOS:3, 4, 6, 5,
2, 7 and 1]. *E. coli* clone PEP1 showed the maximum
15   binding to mAb 2C7 (MFI = 19.81, compared to control
MFI = 4.91), as shown in Figure 3 [SEQ ID NO:1].

   D.   Inhibition ELISA
      Positive *E. coli* clones were grown overnight
in IMC media containing 100 μg/ml ampicillin at 25°C,
20   and then induced to express the peptide fusions for 6
h. Cultures were normalized to the same OD reading
($OD_{600nm} \approx 0.7$), and 1% nonfat dry milk, 150 mM NaCl and
1% α-methyl mannoside were added to block nonspecific
binding. 50 μl-aliquots of each culture were incubated
25   with 50 μl of mAb 2C7 (final concentration 20 ng/ml) at
37°C for 30 min, then 100 μl of the mixtures were
loaded into microtiter plate wells coated with purified
LOS prepared from strain 15253 (80 μg/ml). The wells
were incubated at 37°C for 1 h, then washed. After the
30   wells were washed, bound mAb 2C7 was detected with
anti-mouse IgG conjugated to alkaline phosphatase. A
negative clone that did not bind mAb 2C7 was used as a
control.

PEP1 clones showed the maximum inhibition of
mAb 2C7 binding to LOS (66%) [SEQ ID NO:1]. PEP7,
PEP3, PEP4, PEP2, PEP6, and PEP5 showed respective
decreases in inhibition of binding, as depicted in
5   Figure 4 [SEQ ID NOS:7, 3, 4, 2, 6 and 5]. The
inhibition ELISA results correlated with the flow
cytometric analysis results in that PEP1 also showed
the maximum binding to mAb 2C7. The binding of *E. coli*
cells to mAb 2C7 correlated approximately with
10   decreases in inhibition of mAb 2C7 binding to LOS by
E. coli clones.

II.   Synthetic Peptide Mimic Binding to mAb 2C7

A synthetic peptide (PEP1; IPVLDENGLFAP)
whose sequence corresponds to the consensus sequence
15   "DE_GLF" and includes two cysteine flanking regions
(CGP- and -GPC residues at the - and C- terminus,
respectively) was synthesized (Boston Biomolecules, MA)
to assess specific binding to 2C7 mAb by inhibition
ELISA and to determine whether peptide mimics
20   characterized as thioredoxin-fusion proteins would
retain the antigenicity independent of the fusion
context [SEQ ID NO:10].
The cysteine flanking regions were added to
assess whether antibody binding is affected by
25   cyclization of the peptide mimic. In these peptides
mimics, the cysteine residues allow for the formation
of a disulfide bond between them, resulting in a cyclic
peptide mimic. Such conformationally constrained
peptides may more closely resemble the epitope that
30   they mimic, and therefore may be more immunogenic.
Peptides were diluted in blocking buffer (1%
ovalbumin, 0.05% tween-20, 0.5 M NaCl in PBS) to
produce mixtures of varying concentrations (0.1, 0.5

and 1 mg/ml). 50 μl-aliquots from each of the
concentrations were incubated with 50 μl of mAb 2C7
(stock concentration 2 μg/ml diluted in blocking
buffer) at 37°C for 1 h, then 100 μl of the mixtures
5   were loaded into microtiter plate wells coated with
purified LOS prepared from strain 15253 (80 μg/ml).
The wells were incubated at 37°C for 1 h, then washed.
After the wells were washed, bound mAb 2C7 was detected
with anti-mouse IgG conjugated to alkaline phosphatase.
10   Purified LOS prepared from gonococcal strain 15253 was
used as a positive control. A non-reactive 15-mer
peptide sequence generated by the above described
random peptide library system was used as a negative
control peptide [SEQ ID NO:9].

15   PEP1 inhibited the binding of mAb 2C7 to LOS
in a dose responsive manner (percentage inhibition
equalled 17, 77, and 91% with concentrations of 0.1,
0.5, and 1.0 mg/ml of PEP1, respectively), as shown in
Figure 5. The control 15-mer peptide was synthesized
20   as a cyclic peptide (*CKSNPIHIIKNRRNIPC*) [SEQ ID
NO:9]. This negative control peptide did not inhibit
binding of 2C7 mAb to the purified LOS coated plate.

        Cyclic peptide mimics as described
immediately above may further comprise one or more
25   "tails" for coupling to a second agent, such as an
adjuvant or a carrier protein, by methods known in the
art.

III. Increasing The Immunogenicity Peptide Mimics

        Although small peptides may be immunogenic,
30   several studies have reported that certain small
peptides may lack immunogenicity and result in
ineffective immune responses (particularly humoral

responses) (3, 43). A number of strategies have been used to increase the immunogenicity of small peptides. These include linking the peptide to a carrier protein (54, 28, 54), combining the peptide with an adjuvant

5 (21, 22), using a multiple antigen peptide (MAP) to provide a larger configurational structure that may be more immunogenic (39) and coupling the peptide to a complement protein to enhance the humoral immune response (15).

10 A.   Multiple-antigen Peptide Synthesis

The multiple-antigen peptide (MAP) approach is a technique which associates the peptide mimic with a dendritic matrix of lysine residues (44, 8, 43). Peptides are attached to the amino groups of the lysine

15 scaffold to yield a macromolecule that provides a high density of desired peptide epitopes on the surface of the complex. This approach has been shown to augment the immune response to peptides (39, 40).

A multiple antigen peptide of PEP1 and a

20 control peptide were synthesized (Boston Biomolecules, MA) and binding to mAb 2C7 was assayed by direct and inhibition ELISA.

Solid phase ELISA was performed to assess the binding of mAb 2C7 to multiple antigen peptides. For

25 direct ELISA, Immulon 1 plates were coated overnight with multiple antigen peptides (1 μg/well) and reacted with varying concentration of mAb 2C7. For inhibition ELISA, plates were coated with purified LOS prepared from N. gonorrhoeae strain 15253 (80 μg/ml) at 37°C for

30 3 h. Peptides (linear or MAPs) were diluted in blocking buffer (1% ovalbumin, 0.05% tween-20, 0.5 M NaCl in PBS) to produce mixtures of varying

concentrations. 50 μl-aliquots from each concentration were incubated with 50 μl of mAb 2C7 (stock concentration 0.4 μg/ml diluted in blocking buffer) at 37°C for 1 h, then 100 μl of mixtures were loaded into microtiter plate wells. The wells were incubated at 37°C for 1 h, then washed. After the wells were washed, bound mAb 2C7 was detected with anti-mouse IgG conjugated to alkaline phosphatase. Purified LOS prepared from gonococcal strain 15253 was used as a positive control in inhibition ELISA.

Multiple antigen peptide forms of PEP1 containing four linear PEP1 molecules ("Tetra-MAP1") or eight linear PEP1 molecules ("Octa-MAP1") showed strong binding to mAb 2C7, whereas control MAP showed no binding in direct ELISA, as depicted in Figure 6. Both Tetra- and Octa-MAP1 inhibit mAb 2C7 binding to LOS better than linear PEP1, as depicted in Figure 7. Half maximal inhibition (IC_{50}) for both tetra- and octa-MAP1 was seen at 1.26 μM and 0.23 μM respectively. IC_{50} for linear PEP1 55 μM. This may be due to increased avidity of MAP1 binding to mAb 2C7. Control MAPs showed no significant inhibition.

Immunization with octa-MAP1 induces an IgG anti-LOS antibody response in mice, as shown in Figure 8. The response profile seen in Figure 8(A), in which there is no significant IgG anti-LOS response until the boost at week 3, indicates that the Octa-MAP1 elicited a T-cell dependent immune response in the responding mice. These results demonstrate the promise of a peptide mimic, such as Octa-MAP1, for immunizing humans against *N. gonorrhoeae* infection.

In Figure 8(A), eight mice received a dose of 50 μg of Octa-MAP1 emulsified in Freund's adjuvant on day 0 and again on day 21. Octa-MAP1, which mimics the

2C7 oligosaccharide epitope, induced IgG anti-LOS
antibody in three of the eight mice. IgG anti-LOS
responses in these three mice rose significantly after
the first boost at week 3, peaked at week 7 (the next
5 time measured) and decreased thereafter. Figure 8(B)
shows the positive control experiment in which four
mice were immunized with purified LOS. In these mice,
IgG anti-LOS titers increased minimally after the first
immunization and rose after boosting. All mice in the
10 LOS group showed an anti-LOS antibody response. Four
mice immunized with either Freund's adjuvant (C) or an
unrelated octa-MAP control peptide (D), both negative
controls, elicited weak or no IgG anti-LOS responses.
The mean IgG anti-LOS antibody responses from all
15 immunized mice (from the experiments depicted in Figure
8) are shown in Figure 9 (mean ± SE, including animals
that exhibited no response).

        IgG anti-LOS antibody responses for the
responder mice only (from the experiments depicted in
20 Figure 8) are shown in Figure 10. Antibody response is
defined as IgG anti-LOS (mean ± SE) greater than 0.4
μg/ml (4 fold above baseline IgG anti-LOS levels). At
7 and 10 weeks after primary immunization, responder
mice immunized with Octa-MAP1 elicited IgG anti-LOS
25 antibody levels higher (p < 0.001) than antibody levels
elicited by negative control antigens (Freund's
adjuvant alone or unrelated octa-MAP control peptide).

        IgM anti-LOS antibody responses for responder
mice only (from the experiments depicted in Figure 8)
30 are shown in Figure 11. Mice immunized with Octa-MAP1
that had elicited IgG anti-LOS responses failed to
respond with IgM anti-LOS levels higher than mice
immunized with negative control antigens. Immunization
with LOS (positive control) elicited IgM anti-LOS

antibody levels higher than animals immunized with
either Octa-MAP1 or negative control antigens (Freund's
adjuvant alone or unrelated octa-MAP control peptide).

5   Serum from a mouse immunized with Octa-MAP1
exhibited 2C7-specific complement-mediated bactericidal
activity against *N. gonorrhoeae* strain 15253, as shown
in Figure 12. Depicted in Figure 12 is a graph showing
survival of *N. gonorrhoeae* strain 15253 and its *lgtG*
mutant (2C7 epitope negative) (4) exposed to mouse
10   immune serum (67% final mouse immune serum
concentration by volume) plus added human complement
obtained from normal human donors (17% final human
complement concentration by volume).

Strain 15253 exhibits the 2C7 epitope.
15   Strain 15253 *lgtG* contains a disrupted allele of
lipooligosaccharide (LOS) glycosyl transferase G, which
transfers glucose (via an α linkage) onto heptose-2 in
the core of LOS (4). The disruption of the *lgtG* locus
results in the loss of 2C7 epitope expression.

20   A standard bactericidal assay was performed
to assess complement-mediated bactericidal activity in
mouse sera (11). In this assay, mouse serum (67% final
volume) (from various mice immunized or not as
described below) was incubated with approximately 2.5 X
25   10^3 bacteria suspended in Morse A media (33) in the
presence of human complement (17% final volume). The
reaction mixture was then shaken continuously at 37°C
for 30 minutes. Aliquots of the reaction mixture were
then inoculated onto chocolate agar plates at time 0
30   and 30 minutes. Survival was expressed as the percent
increase in colonies on the plate at 30 minutes,
compared to those on the plate at 0 minutes. Greater
than 100% survival in the assay indicates growth during
the 30-minute incubation period.

mAb 2C7 was used as a control, as it kills *N. gonorrhoeae* strain 15253 with added complement, but does not kill the 15253 *lgtG* mutant strain. As shown in Figure 12(A), mAb 2C7 possesses bactericidal

5   activity against 2C7 epitope-bearing gonococci. 25 µg/ml of mAb 2C7 (100 µl in 150 µl of total volume of reaction mixture) mediated 100% killing of strain 15253, and no killing of strain 15253 *lgtG*.

Serum taken from a single mouse immunized

10 with Octa-MAP1, containing 5.05 µg/ml of IgG anti-LOS antibody pooled from bleeds taken between weeks 7-11, showed 92% killing (8% survival) of strain 15253 whereas strain 15253 *lgtG* survived fully, as depicted in Figure 12(C).

15 Normal mouse serum representing a pool of 20 mouse sera with a mean concentration of IgG anti-LOS antibody of 0.1 µg/ml failed to kill either strain, as shown in Figure 12(B). Control mouse serum without complement showed 116.1% ± 4.7% survival (no killing)

20 for strain 15253, and 123.1% ± 3.5% survival (no killing) for the *lgtG* mutant of 15253. The complement source without antibody exhibited 137.9% ± 1.0% survival (no killing) for strain 15253, and 132.5% ±14.3% survival (no killing) for the *lgtG* mutant of

25 15253.

Serum taken from a single mouse immunized with LOS (containing 21.98 µg/ml of IgG anti-LOS antibodies, pooled from bleeds taken between weeks 7-11) effected no killing of strain 15253 (179% survival)

30 and strain 15253 *lgtG* (133% survival), as shown in Figure 12(D). Serum taken from single mice immunized with Freund's adjuvant alone or unrelated Octa-MAP

control peptide, as negative control antigens, did not kill either strain, as shown in Figures 12(E) and 12(F) respectively.

IgG anti-LOS antisera obtained from mice
5 immunized with Octa-MAP1 exhibited concentration-dependent killing of *N. gonorrhoeae* strain 15253, as shown in Figure 13.

Figure 13 shows a plot of IgG anti-LOS antibody concentration versus killing of *N. gonorrhoeae*
10 strain 15253. When IgG anti-LOS antisera levels from each of three mice immunized with Octa-MAP1 were plotted against bacterial killing, a dose-response profile resulted (mouse sera containing 1.38, 2.50 and 5.05 μg/ml of anti-LOS antibodies showed 31, 74 and 92
15 % killing, respectively, of strain 15253). Killing by mAb 2C7 was also shown at 5 separate LOS antibody concentrations as a positive control.

B.   Coupling A Peptide Mimic
To Complement Protein C3d

20   It is expected that the immunogenicity of peptide mimics of gonococcal epitopes, such as Octa-MAP1 described herein, can be further enhanced through coupling with complement factor C3d.

Numerous studies have demonstrated an important
25 role of complement protein C3 in the induction of humoral immune responses (1, 5, 14, 17, 25, 32, 34 and 35). C3-depleted mice show diminished antibody responses to T-cell dependent protein antigen, such as keyhole limpet haemocyanin ("KLH") (34, 35).
30 Complement receptor 1-(CR1 or CD35) and complement receptor 2-(CR2 or CD21) deficient mice have an impaired T-cell dependent antibody response (1, 14,

32). It has further been shown that C3d covalently
linked to hen egg lysozyme ("HEL") resulted in an
enhanced antibody response to the HEL antigen (15).
Mice immunized with a fusion protein that consisted of
5 three copies of C3d and 1 copy of HEL elicited a
10,000-fold increase in anti-HEL antibody response,
compared to antibody responses in mice immunized with
HEL alone. Anti-HEL antibody responses induced by the
fusion protein were approximately 100- fold higher than
10 those induced by HEL emulsified in Freund's adjuvant.

Octa-MAP1 can be coupled to C3d by cloning an
octa-MAP1 DNA sequence into a C3d fusion protein
cassette and transforming this construct into an
expression system. The OctaMAP1-C3d fusion protein can
15 then be expressed, purified and used as an immunogen.
Alternatively, the OctaMAP1-C3d gene fusion, in the
form of DNA, can be used as a DNA vaccine according to
methods known in the art.

A hybridoma producing anti-idiotypic
20 antibodies exhibiting immunological reactivity similar
to the peptide mimics of the instant invention is
exemplified by a cell culture deposited in the ATCC
(10801 University Boulevard, Manassas, Va. 20110-2209
U.S.A.) on March 26, 1993 and assigned ATCC accession
25 number HB 11311.

Hybridoma 2C7 secreting the mAb 2C7
exhibiting immunological reactivity similar to the
peptide mimics of the instant invention is exemplified
by a cell culture designated as 2C7 and deposited in
30 the ATCC on March 9, 1995. This culture was assigned
ATCC accession number HB-11859.

While we have hereinbefore described a number
of embodiments of this invention, it is apparent that
our basic constructions can be altered to provide other

embodiments which utilize the processes and
compositions of this invention. Therefore, it will be
appreciated that the scope of this invention is to be
defined by the claims appended hereto rather than by
5   the specific embodiments which have been presented
hereinbefore by way of example.

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